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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.	
10/549,340	09/13/2005	Takayuki Abe	114/75034 6716		
23432 COOPER & D	23432 7590 01/15/2008 COOPER & DUNHAM, LLP			EXAMINER	
1185 AVENUE OF THE AMERICAS			CWERN, JONATHAN		
NEW YORK, NY 10036			ART UNIT	PAPER NUMBER	
			3737		
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Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

	Application No.	Applicant(s)
	10/549,340	ABE ET AL.
Office Action Summary	Examiner	Art Unit
	Jonathan G. Cwern	3737
The MAILING DATE of this communication app Period for Reply	ears on the cover sheet with the c	orrespondence address
A SHORTENED STATUTORY PERIOD FOR REPLY WHICHEVER IS LONGER, FROM THE MAILING DA  - Extensions of time may be available under the provisions of 37 CFR 1.13 after SIX (6) MONTHS from the mailing date of this communication.  - If NO period for reply is specified above, the maximum statutory period w  - Failure to reply within the set or extended period for reply will, by statute, Any reply received by the Office later than three months after the mailing earned patent term adjustment. See 37 CFR 1.704(b).	ATE OF THIS COMMUNICATION 36(a). In no event, however, may a reply be tim vill apply and will expire SIX (6) MONTHS from cause the application to become AB ANDONE	I.  nely filed  the mailing date of this communication.  D (35 U.S.C. § 133).
Status	•	
1)⊠ Responsive to communication(s) filed on <u>09 Note</u> 2a)⊠ This action is <b>FINAL</b> . 2b)□ This     3)□ Since this application is in condition for alloward closed in accordance with the practice under E	action is non-final. nce except for formal matters, pro	
Disposition of Claims	<b>,</b>	
4) Claim(s) 1-3 and 5-23 is/are pending in the app 4a) Of the above claim(s) is/are withdraw 5) Claim(s) is/are allowed. 6) Claim(s) 1-3 and 5-23 is/are rejected. 7) Claim(s) is/are objected to. 8) Claim(s) are subject to restriction and/or Application Papers	vn from consideration.	
9) The specification is objected to by the Examine 10) The drawing(s) filed on is/are: a) access Applicant may not request that any objection to the confidence of Replacement drawing sheet(s) including the correction 11) The oath or declaration is objected to by the Examine 11.	epted or b) objected to by the Edrawing(s) be held in abeyance. See on is required if the drawing(s) is obj	e 37 CFR 1.85(a). ected to: See 37 CFR 1.121(d).
Priority under 35 U.S.C. § 119		
12) Acknowledgment is made of a claim for foreign  a) All b) Some * c) None of:  1. Certified copies of the priority documents  2. Certified copies of the priority documents  3. Copies of the certified copies of the prior  application from the International Bureau  * See the attached detailed Office action for a list of	s have been received. s have been received in Application ity documents have been received (PCT Rule 17.2(a)).	on No ed in this National Stage
Attachment(s)  1) Notice of References Cited (PTO-892)  2) Notice of Draftsperson's Patent Drawing Review (PTO-948)  3) Information Disclosure Statement(s) (PTO/SB/08)  Paper No(s)/Mail Date	4)  Interview Summary Paper No(s)/Mail Da 5)  Notice of Informal P 6) Other:	te

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## **DETAILED ACTION**

## Claim Rejections - 35 USC § 103

The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negatived by the manner in which the invention was made.

This application currently names joint inventors. In considering patentability of the claims under 35 U.S.C. 103(a), the examiner presumes that the subject matter of the various claims was commonly owned at the time any inventions covered therein were made absent any evidence to the contrary. Applicant is advised of the obligation under 37 CFR 1.56 to point out the inventor and invention dates of each claim that was not commonly owned at the time a later invention was made in order for the examiner to consider the applicability of 35 U.S.C. 103(c) and potential 35 U.S.C. 102(e), (f) or (g) prior art under 35 U.S.C. 103(a).

Claims 1-3, and 5-23 are rejected under 35 U.S.C. 103(a) as being unpatentable over Mistretta et al. (US 5713358) in view of Ho et al. (US 2002/0087069).

Mistretta shows, means for dividing k space into high repetitive-frequency areas containing the origin and low repetitive-frequency areas not containing the origin (column 7, line 65-column 8, line 65), signal processing means for reconstructing an image using the k space data (column 8), a display (column 5, lines 20-40), labeling the

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time period of the sampled data so that it can be used to reconstruct the image (column 8, lines 25-40), reconstructing the image using the high frequency area measurement and a low frequency area measurement which are close in time (column 8 line 45-column 9, line 15); a lower frequency area is measured immediately after a high frequency area (column 8); all of k space is sampled (column 8); k space data is data of concentration information for a contrast medium injected into a blood vessel (column 10); k space comprises a slice encode direction, a phase encode direction, a readout direction, and k space is divided by a plane parallel to the readout direction (columns 7 and 8); projection processing on a two-dimensional plane after three-dimensional reconstruction (column 10, lines 30-50);

Mistretta fails to show, comparing a time phase evaluation value with a predetermined threshold value; controlling the measurement sequence so that the high frequency area contains the time phase; predicting the timing from the time change of the time phase evaluation value, controlling the measurement sequence based on the timing predicted; determining the time phase after the measurement repetitions; the time phase value is a peak value of the k space data; the time phase value is a value of data which has been fourier transformed; the threshold value is at least 1.8 times the baseline value of the time phase evaluation value; the threshold value is at least 80% of a maximum value of the time phase evaluation value; display has means for setting the threshold value, designating the time phase, selecting each measurement area; displaying the time phase evaluation value in a time series, a signal intensity change curve, a measurement sequence of the measurement areas, differing display of each

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measurement area selected from those not selected; and a time phase value is a value in which an artery is emphasized by contrast agent.

Ho teaches, comparing a time phase evaluation value with a predetermined threshold value ([0037]-[0041]); controlling the measurement sequence so that the high frequency area contains the time phase (high frequency areas are also acquired with the same steps, [0045]-[0048]); predicting the timing from the time change of the time phase evaluation value ([0037]-[0041]), controlling the measurement sequence based on the timing predicted ([0037]-[0041]); determining the time phase after the measurement repetitions ([0037]-[0041]); the time phase value is a peak value of the k space data (the value is when the contrast agent enters the region of interest, the presence of the contrast agent will increase the value so that it is at its highest point, a peak [0037]-[0041]); the time phase value is a value of data which has been fourier transformed ([0029]); the threshold value is at least 1.8 times the baseline value of the time phase evaluation value (the threshold is preselected by the user, and so it can be selected to be any value, [0040]); the threshold value is at least 80% of a maximum value of the time phase evaluation value (the threshold is preselected by the user, and so it can be selected to be any value. [0040]); display has means for setting the threshold value, designating the time phase, and selecting each measurement area (the system is operator controlled, by any of various input devices, so that the operator could control any of these variables, [0026]); displaying the time phase evaluation value in a time series, a signal intensity change curve, a measurement sequence of the measurement areas, differing display of each measurement area selected from those

not selected (a display will be capable of displaying any of these images, [0026]); and a time phase value is a value in which an artery is emphasized by contrast agent ([0037]).

Mistretta mentions labeling the time periods during which the k space data is obtained, and using the time periods to reconstruct the data at a later time. However, he does not go into further detail on how this is accomplished. Ho describes in detail how a threshold can be used to mark the time periods during which data is acquired. It would have been obvious to one of ordinary skill in the art, at the time the invention was made, to have used a threshold to mark the time periods during which data is acquired, in the system of Mistretta, with the motivation that marking the time of the data acquisition will aid in reconstructing the high and low frequency areas, so that image will be easier for the physician to evaluate.

## Response to Arguments

Applicant's arguments filed 11/9/07 have been fully considered but they are not persuasive.

In regards to applicant's argument that Ho does not involve changing the predetermined sequence, while repeating measurement of the measurement areas, of each of said measurement areas in such a manner that a measurement period of said high repetitive frequency measurement area contains said time phase, examiner respectfully disagrees. First, by changing the start timing of the measurement sequence, Ho does in fact meet the claim limitation which states: "said measurement control means changes the predetermined measurement sequence of each of said

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measurement areas in such a manner that a measurement period of said high repetitive-frequency measurements contains said time phase". The arterial phase is specifically mentioned for example in paragraph [0051], the arterial phase being a measurement period during which high repetitive-frequency measurements contain the time phase. In addition, in paragraph [0055] Ho describes moving the patient table so as to chase the bolus. The acquisition time is changed on-the-fly based on the passage of the contrast bolus. Therefore the system will change the measurement sequence based on the speed of the bolus, tracking the bolus during the arterial phase.

In regards to applicant's arguments that the cited art does not disclose selecting the high and low areas as an image reconstruction set, examiner respectfully disagrees. As stated in the previous non-final rejection, Mistretta et al. clearly show that an image is reconstructed using data from a central k-space region and temporally adjacent data from the surrounding peripheral k-space regions (column 8, line 45-column 9, line 15). One method described is using the data acquired from peripheral regions closest in time to the acquisition of the central k-space region. Each image frame data set depicts the subject at a particular time during the study. Ho et al. also disclose combining high and low spatial frequency data sets to reconstruct and image of the arterial vasculature ([0053]). In addition, in applicant's specification, on pages 18-19, applicant submits that this is a known technology disclosed in JP-A-2002-177240, dividing k-space into high and low frequency areas and then using the data for image reconstruction.

Therefore the previous rejection dated 8/9/07 is upheld and repeated above, including new claims 22 and 23.

## Conclusion

The prior art made of record and not relied upon is considered pertinent to applicant's disclosure.

Prince (US 5590654) discloses a technique and apparatus for monitoring and detecting the arrival of a contrast agent in a region of interest, which may be employed to facilitate synchronization between collecting the central portion of k-space image data with the arterial phase of contrast enhancement.

Applicant's amendment necessitated the new ground(s) of rejection presented in this Office action. Accordingly, **THIS ACTION IS MADE FINAL**. See MPEP § 706.07(a). Applicant is reminded of the extension of time policy as set forth in 37 CFR 1.136(a).

A shortened statutory period for reply to this final action is set to expire THREE MONTHS from the mailing date of this action. In the event a first reply is filed within TWO MONTHS of the mailing date of this final action and the advisory action is not mailed until after the end of the THREE-MONTH shortened statutory period, then the shortened statutory period will expire on the date the advisory action is mailed, and any extension fee pursuant to 37 CFR 1.136(a) will be calculated from the mailing date of the advisory action. In no event, however, will the statutory period for reply expire later than SIX MONTHS from the date of this final action.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Jonathan G. Cwern whose telephone number is 571-270-1560. The examiner can normally be reached on Monday through Friday 9:30AM - 6:00PM EST.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Brian Casler can be reached on 571-272-4956. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free). If you would like assistance from a USPTO Customer Service Representative or access to the automated information system, call 800-786-9199 (IN USA OR CANADA) or 571-272-1000.

JC

/Ruth S. Smith/ Ruth S. Smith Primary Examiner Art Unit 3737